In the claims:

Please amend claims 8, 9, and 14 and insert new claims 16 and 17 as shown below.

1. (Original) A pharmaceutical composition for the treatment of wounds, comprising a pharmaceutically effective amount of blood plasma as an active agent.

2. (Original) The pharmaceutical composition according to claim 1, wherein pH is in the range of 3.5 to 6.6

3. (Original) The pharmaceutical composition according to claim 1, wherein the active agent is derived from livestock.

4. (Original) The pharmaceutical composition according to claim 1, wherein it is topically administered.

5. (Original) The pharmaceutical composition according to claim 1, in the form of creams, ointments, gels, liquids or patched.

6. (Original) The pharmaceutical composition according to claim 1, wherein the wounds include contusion or bruise, non-healing traumatic wounds, the disruption by irradiation, abrasion, bone gangrene, laceration, avulsion, penetrated wound, gun shot wound, cutting, burn, cold sores, cutaneous ulcers, xeroderma, skin kefatosis, break, rupture, dermatitis, pain by dermatophyte, wounds by surgery or by vascular disorder, corneal wounds, pressure sore, bed sore, certain conditions associated with diabetes such as diabetic cutaneous disorder and with

poor circulation, chronic ulcers, suture site caused by plastic surgery, spinal traumatic wounds, gynecological wounds, chemical wounds and acne.

7. (Currently amended) The pharmaceutical composition according to ~~claims~~ claim 1 ~~or 6~~, which is used at an amount of from 0.01 to 0.1 g/cm^2 in the treatment of full thickness defect wounds.

8. (Original) A pharmaceutical composition for the treatment of wounds, comprising a pharmaceutically effective amount of blood serum as an active agent.

9. (Original) The pharmaceutical composition according to claim 8, wherein pH is in the range of 3.5 to 6.6

10. (Original) The pharmaceutical composition according to claim 8, wherein the active agent is derived from livestock.

11. (Original) The pharmaceutical composition according to claim 8, wherein it is topically administered.

12. (Original) The pharmaceutical composition according to claim 8, in the form of creams, ointments, gels, liquids or patched.

13. (Original) The pharmaceutical composition according to claim 8, wherein the wounds include contusion or bruise, non-healing traumatic wounds, the disruption by irradiation, abrasion, bone gangrene, laceration, avulsion, penetrated wound, gun shot wound, cutting, burn, cold sores, cutaneous ulcers, xeroderma, skin kefatosis, break, rupture, dermatitis, pain by dermatophyte, wounds by surgery or by vascular disorder, corneal wounds, pressure sore, bed sore, certain conditions associated with diabetes such as diabetic cutaneous disorder and with poor circulation, chronic ulcers, suture site caused by plastic surgery, spinal traumatic wounds, gynecological wounds, chemical wounds and acne.

14. (Currently amended) The pharmaceutical composition according to ~~claims~~ claim 8 ~~or 13~~, which is used at an amount of from 0.01 to 0.1 g/cm^2 in the treatment of full thickness defect wounds.

15. (New) The pharmaceutical composition according to claim 6, which is used at an amount of from 0.01 to 0.1 g/cm^2 in the treatment of full thickness defect wounds.

16. (New) The pharmaceutical composition according to claim 13, which is used at an amount of from 0.01 to 0.1 g/cm^2 in the treatment of full thickness defect wounds.